APPENDIX N BULK MILK TANKER SCREENING TEST FORM

GENERAL REQUIREMENTS

(Unless otherwise stated all tolerances ±5%)

1.	Wor	k Area	
	a.	Ample working space and utilities	
	b.	Clean well ventilated, test kit used in temperature range specified by manufacturer, reasonably free from dust and drafts	
	c.	Adequate lighting, [NCIMS CERTIFIED LABORATORIES and CERTIFIED INDUSTRY SUPERVISORS, > 50 foot-candles at working surface (pref 100)]	
2.	Sto	rage Space	
	a.	Cabinets, drawers, and shelves adequate	
	b.	Areas neat, clean and orderly	
3.	 b. Clean well ventilated, test kit used in temperature range specified by manufacturer, reasonably free from dust and drafts		
	a.	Thermometer traceable to NIST Certified thermometer	
	b.	Range of thermometers appropriate for designated use	
	C.	CERTIFIED LABORATORIES and CERTIFIED INDUSTRY	
	d.	traceable thermometer annually (including	
		1. Accurate to ±1C	
		date, identification, temperature checked and	
	e.	Dial thermometers not permitted	

1 .	Ref:	rigeration	
	a.	Size adequate for workload	
	b.	Maintains samples at 0-4.4C	
	c.	Reagents stored as per manufacturer instructions	
	d.	Not used to store food or drink for consumption	
	e.	Record temperature daily from 2 thermometers with bulbs submerged in liquid, placed on upper and lower shelves of use [NCIMS CERTIFIED LABORATORIES and CERTIFIED INDUSTRY SUPERVISORS, AM and PM]	
	f.	NCIMS CERTIFIED LABORATORIES and CERTIFIED INDUSTRY SUPERVISORS, dedicated for milk work only	
5.	Fre	ezer _	
	a.	Size adequate for workload	
	b.	Maintains -15C or below	
	c.	Not used to store food or drink for consumption	
	d.	Record temperature daily	
	e.	NCIMS CERTIFIED LABORATORIES and CERTIFIED INDUSTRY SUPERVISORS, dedicated for milk work only, NO PATHOGENS STORED	
6.	Bal	ance, electronic (if necessary)	
	a.	Weight capability appropriate for intended use	
	b.	Accurate to 0.01g for preparations of positive controls	
	c.	Appropriate sensitivity for calibration of pipetting devices within a tolerance of ±5% (0.001g sensitivity appropriate in most instances)	
		1. Pipetting devices calibrated on-site	
		2. Pipetting devices calibrated at another location	
	d.	Checked monthly with Class S or S1, or equivalent ASTM 1, 2 or 3, weights (Appendix N drug testing only laboratories may check quarterly)	
	e.	Checked annually by a qualified service representative	

(APPN/GENREQ-2-Rev. 9/00)

	f.	Records maintained	
7.	[Re	ettors, calibrated, fixed volume type only quired for NCIMS Certified Laboratories and tified Industry Supervisors]	
	a.	Calibrate with ten (10) consecutive weighings quarterly (using separate tip for each weighing)	
	b.	Average of all 10 weighings must be $\pm 5\%$ of specified delivery volume (≤ 1.0 mL by weight, > 1.0 mL by volume using class A graduated cylinder), records maintained	
	c.	Etched with identification and tagged with date calibrated	
	d.	Appropriate tips for pipettor(s) used	
8.		onized Water or Equivalent, or as specified manufacturer	
		SAMPLES	
9.	Sam	ple Requirements	
	a.	Ascertain temperature of bulk milk tanker	
	b.	Secure a representative sample for drug residue testing	
	c.	Prevent contamination with disinfectants from hands or other sources	
	d.	Samples tested and confirmation completed (when necessary) within 72 hours of initial collection	
		ADDITIONAL REQUIREMENTS FOR NCIMS CERTIFIED LABORATORIES AND CERTIFIED INDUSTRY SUPERVISORS	
	e.	Record time, date and temperature of samples as received and tested	
	f.	Determine sample temperature by inserting pre-cooled thermometer (pre-cooling of electronic/digital thermometer probes is not necessary)into temperature control (TC), if no TC, aliquot samples for testing and measure temperature using one of the producer samples	
	g.	Do not accept producer samples (about ¾ full) that are over filled	

	h.	If raw milk exceeds 4.4C on receipt do not test (samples may be received at 7C if time of receipt is ≤ 3 hours from collection and arrival temperature is equal to or less than temperature of collection)	
		PERFORMANCE TESTING	
10.	Per	formance Testing	
	a.	Run a positive and negative control with each new lot of kits, must give appropriate results, records maintained	
	b.	Run a negative and positive control DAILY (on days testing), at each test site, must give appropriate results, if not, re-run controls (may be necessary to prepare new controls), if problem persists discontinue testing, contact State regulatory and seek technical assistance, records maintained	
	c.	If more than one analyst performs analysis have different analyst run performance check on rotational basis	
		CONFIRMATION OF PRESUMPTIVE POSITIVE SAMPLES	
11.		firmation of Presumptive Positive Samples [Must ply with current edition of M-a-86]	
	a.	The SAME sample is re-tested in DUPLICATE along with a positive and negative control using fixed volume pipettors	
	b.	Positive and negative controls give the appropriate result(s), report sample results	
	C.	If positive and/or negative control do not give appropriate results, re-run controls and samples, if problem persists contact state regulatory and seek technical assistance	
	d.	Maintain copies of all confirmation reports	
		REPORTING AND RECORDS	
12.	Rep	orting and Records	
	a.	Report as Positive (+) for beta-lactam, specific drug or inhibitor (when a non-specific microbial inhibitor test used without beta-lactamase) when demonstrated	
	b.	Report as Not Found (NF) when demonstrated	
(AP	APPN/GENREQ-4-Rev. 9/00)		

	C.	Record test performed, interpretation of unknowns (samples) and controls	
	d.	Records, including all printouts, maintained for 6 months [NCIMS CERTIFIED LABORATORIES and CERTIFIED INDUSTRY SUPERVISORS, for 2 years]	
		MISCELLANEOUS	
13.	Mis	cellaneous	
	a.	Material safety data sheets (MSDS) on file	
	b.	Current, applicable survey forms available in laboratory	
	c.	Positive Report Forms available with instructions	
	d.	Personnel adequately trained	
	e.	Required split/check sample participation	